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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
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09/763,793

02/27/2001

Hiromasa Miyaji

766.46

3687

5514

7590

10/31/2008

FITZPATRICK CELLA HARPER & SCINTO
30 ROCKEFELLER PLAZA
NEW YORK, NY 10112

EXAMINER

SHAFFER, SHULAMITH H

ART UNIT

PAPER NUMBER

1647

MAIL DATE

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10/31/2008

PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary	Application No. 09/763,793	Applicant(s) MIYAJI ET AL.	
	Examiner SHULAMITH H. SHAFER	Art Unit 1647	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 01 August 2008.
- 2a) ☒ This action is **FINAL**. 2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1-12 and 46-48 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 1-12 and 46-48 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☒ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☒ All b) ☐ Some * c) ☐ None of:
1. ☒ Certified copies of the priority documents have been received.
2. ☐ Certified copies of the priority documents have been received in Application No. _____.
3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- | | |
|--|---|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892) | 4) <input type="checkbox"/> Interview Summary (PTO-413) |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948) | Paper No(s)/Mail Date. _____ |
| 3) <input checked="" type="checkbox"/> Information Disclosure Statement(s) (PTO/SB/08) | 5) <input type="checkbox"/> Notice of Informal Patent Application |
| Paper No(s)/Mail Date <u>8/1/08, 9/2/08</u> . | 6) <input type="checkbox"/> Other: _____ |

Detailed Action

Status of Application, Amendments, And/Or Claims:

A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on 1 August 2008 has been entered.

Claims 1-12 and 46-48 are pending in the instant application and are under consideration. These are the same claims that were submitted on 27 March 2008 and entered as an after final amendment.

Information Disclosure Statement:

The Information Disclosure statements (IDS) submitted on the 1 August 2008 and 2 September 2008 have been considered. The signed copies are attached.

Priority:

Acknowledgment is made of applicants' claim for foreign priority based on an application filed in Japan on 27 of August 1998. A certified copy of the Japan 10/241248 application as required by 35 U.S.C. 119(b). Applicants have provided a certified translation of 10/241248 with submission of 1 August 2008; therefore, Applicants have perfected priority claim to 27 August 1998, the date of filing of Japan 10/241248.

Withdrawn Objections/Rejections

35 U.S.C. § 112, First Paragraph:

The rejection of claims 2, 5, 6, 8-12, 29, 42, 46 and 48 under 35 U.S.C. 112, first paragraph (scope of enablement), is withdrawn in view of applicants amendments to the claims.

The rejection of Claims 2, 5, 6, 8-12, 29, 42, 46 and 47 under 35 U.S.C. 112, first paragraph, as failing to comply with the written description requirement is withdrawn in view of applicants' amendment to the claims.

Maintained Rejections

35 U.S.C. §§ 101 and 112, First Paragraph:

35 U.S.C. § 101 reads as follows:

Whoever invents or discovers any new and useful process, machine, manufacture, or composition of matter, or any new and useful improvement thereof, may obtain a patent therefor, subject to the conditions and requirements of this title.

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

The rejection of Claim(s) 1-12 and 46-48 under 35 U.S.C. 101 because the claimed invention is not supported by either a credible, substantial or specific asserted utility or a well established utility is maintained for reasons of record and reasons set forth below.

Applicants traverse the rejection (After final submission of 27 March 2008 and Remarks of 1 August 2008).

The reasons for the traversal are:

a. the polypeptide of SEQ ID NO:1 is a member of the nucleoside transporter system. Dipyridamole, a compound known to inhibit the uptake of adenosine, enhances bronchospasm in an asthmatic patient (Crimi et al. 1988 Allergy 43:179-83, submitted with after-final amendment). Thus, one would conclude that bronchospasm is suppressed by accelerating uptake of adenosine; therefore, when a DNA encoding SEQ ID NO:1 is expressed in the lung of asthmatic patients, bronchospasm in the patient is treated (submission of 27 March 2008, page 7, last paragraph, bridging page 8, 1st paragraph).

b. dipyridamole, as an inhibitor of equilibrative nucleoside transporters, including the one described in the instant invention, inhibits the decrease of extracellular adenosine concentration by inhibiting uptake of adenosine. The transporters take in and eliminate extracellular adenosine. Thus, when transporters are inhibited, more adenosine is available in the extracellular space to activate the adenosine receptor on the cell membrane and cause bronchoconstriction (Remarks of 1 August 2008, page 2, numbered paragraphs 1-3).

c. References provided with applicants submission provide evidence that concentration of adenosine is elevated in the lung of asthmatic and adenosine-induced bronchoconstriction is mediated by A(1) receptor on the cell membrane.

Response to Arguments

Applicant's arguments have been fully considered but are not found to be persuasive for the following reasons:

Applicants have sought to establish the following fact pattern:

1. Adenosine, present in elevated levels in the lungs of asthmatic patients, binds to the A(1) receptor, and elicits bronchoconstriction (Hua et al. 2007. American Journal of Physiology-Lung, Cellular and Molecular Physiology. 293:L25-32 and Brown et al. 2008. Eur. Resp J. 31:311-9, abstracts of both references submitted on 1 August 2008).
2. Nucleoside transporters, including the polypeptide of the instant invention, transport adenosine into the cell, thereby reducing the amount of adenosine available to

Art Unit: 1647

activate the A(1) receptor; thus these polypeptides play a role in reducing bronchospasms.

3. Therefore, the polypeptide of the instant invention and its encoding DNA would be useful in treating bronchospasm in the asthmatic patient.

In summary, Applicants assert, in above arguments (a, b, 2 and 3 above), that the polypeptide of the instant invention and DNA encoding said polypeptide have utility in treating bronchospasm in an asthmatic patient, for which there is no support in the specification of the instant invention.

Applicants are reminded that a specific or substantial asserted utility or a well established utility must be presented at the time of filing. The specification has not asserted a specific and substantial utility nor is there a well established utility for the claimed invention because the specification and/or the art fail to establish a connection between the polypeptide of SEQ ID NO:1 structure, expression or activity or changes in structure, expression or activity and any specific disease state nor has this been established for the encoding DNA (SEQ ID NO:2). Applicants assert, in the specification, that the polypeptide of the instant invention may be used as "a preventive agent or a therapeutic agent for ischemic heart disease, cerebral disorder at the time of stroke, immune response accompanied by organ transplantation, malignant tumor, nephritis, pancreatitis or hypertension....Its applications as an analgesic, an antiplatelet agent, an agent for increasing activity of an antiviral agent or a malignant tumor treating agent and an agent for reducing side effects at the time of chemotherapy can also be expected" (page 63, last paragraph, bridging page 64, 1st paragraph). There is no assertion that the polypeptide of the instant invention (or its encoding DNA) may be used to treat symptoms in the asthmatic patient.; thus there is no support in the specification for this utility.

In response to a: Applicants argue, without supporting evidence, that when a DNA encoding SEQ ID NO:1 is expressed in the lung of asthmatic patients, bronchospasm in the patient is treated by gene therapy. The specification does not

Art Unit: 1647

contemplate any gene therapy methods or protocols and does not support this assertion of utility.

In response to b: One of skill in the art would be unable to predict that nucleoside transporters, such as the polypeptide of the instant invention would have a therapeutic role in treatment of asthma and related bronchospasms. The fact that dipyridamole, an inhibitor of equilibrative nucleoside transporters, enhances bronchospasm in an asthmatic patient does not provide evidence that stimulating nucleoside transporters or increasing the level of nucleoside transporter protein in the cell would inhibit bronchospasms. Contrary to applicants' assertion above, the art teaches that equilibrative transporters can move adenosine bidirectionally across plasma membranes by facilitated diffusion. Adenosine formed intracellularly can be released by bidirectional nucleoside transport processes to activate cell surface receptors (Borgland et al. 1998, Europ. J of Pharm. 346:339-344, abstract and page 339, 2nd column, 1st paragraph). Thus, the presence of additional nucleoside transporter polypeptides on the surface of bronchial cells might stimulate bronchospasms, by transporting adenosine to the extracellular spaces.

Further research would be required to ascertain the function of SEQ ID NO:1, and to identify a disease with which this polypeptide is associated. Thus, the instant application is an invitation to the skilled artisan to experiment as to the function of the polypeptide of the instant invention and to determine if there is any nexus between said polypeptide any disease or pathological condition.

Utility must be in readily available form. In *Brenner v. Manson*, 148 U.S.P.Q. 689 (Sup. Ct., 1966), a process of producing a novel compound that was structurally analogous to other compounds which were known to possess anti-cancer activity was alleged to be useful because the compound produced thereby was potentially useful as an anti-tumor agent in the absence of evidence supporting this utility. The court expressed the opinion that all chemical compounds are "useful" to the chemical arts when this term is given its broadest interpretation. However, the court held that this

Art Unit: 1647

broad interpretation was not the intended definition of "useful" as it appears in 35 U.S.C. § 101, which requires that an invention must have either an immediately obvious or fully disclosed "real world" utility. The instant claims are drawn to a polynucleotide encoding a protein which has undetermined function or biological significance. Until some actual and specific activity can be attributed to the protein identified in the specification as SEQ ID NO:1 or the polynucleotides encoding it (SEQ ID NO:2) the claimed invention is incomplete.

Since the polypeptide of SEQ ID NO:1, or its encoding nucleic acid molecule (SEQ ID NO:2) are not supported by a specific and substantial utility, or a well-established utility, then expression vectors, and transformants comprising the nucleic acids also do not possess utility.

The rejections of Claims 1-12, and 46-48 under 35 U.S.C. 112, first paragraph are maintained for reasons of record. Specifically, since the claimed invention is not supported by either a specific or substantial asserted utility or a well established utility for the reasons of record and those set forth above, one skilled in the art clearly would not know how to use the claimed invention.

Art made of record:

The following art is made of record and not relied upon is considered pertinent to applicant's disclosure. Baker et al. (WO 200012708) teach a polynucleotide (SEQ ID NO:78), encoding a PRO 1380 polypeptide, which has 99.2% identity to SEQ ID NO:2 of the instant invention (See alignment below). However, the reference claims priority to provisional filed 3 November 1998, which is after the perfected priority date of the instant application (27 August 1998)

Human PRO1380 (UNQ717) cDNA sequence SEQ ID NO:78.
WO200012708-A2.
09-MAR-2000.

Art Unit: 1647

Baker K, Goddard A, Gurney AL, Smith V, Watanabe CK, Wood WI;
Sequence 2243 BP; 463 A; 701 C; 572 G; 507 T; 0 U; 0 Other;

Query Match 99.2%; Score 2223; DB 3; Length 2243;
Best Local Similarity 99.8%; Pred. No. 0;
Matches 2226; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

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Qy      1 CGGCGGCGTGGCGCAGCGGCGACATGGCCGTTGTCTCAGAGGACGACTTTCAGCACAGTT 60
      |||
Db     13 CGGCGGCGTGGCGCAGCGGCGACATGGCCGTTGTCTCAGAGGACGACTTTCAGCACAGTT 72

Qy     61 CAAACTCCACCTACGGAACCACAAGCAGCAGTCTCCGAGCTGACCAGGAGGCACTGCTTG 120
      |||
Db     73 CAAACTCCACCTACGGAACCACAAGCAGCAGTCTCCGAGCTGACCAGGAGGCACTGCTTG 132

Qy    121 AGAAGCTGTGTGGACCGCCCGCCCTGGCCTGCAGAGGCCCGAGGACCGCTTCTGTGGCA 180
      |||
Db   133 AGAAGCTGTGTGGACCGCCCGCCCTGGCCTGCAGAGGCCCGAGGACCGCTTCTGTGGCA 192

Qy    181 CATAATCATCTTCTTTCAGCCTGGGCATTGGCAGTCTACTGCCATGGAACCTCTTTATCA 240
      |||
Db   193 CATAATCATCTTCTTTCAGCCTGGGCATTGGCAGTCTACTGCCATGGAACCTCTTTATCA 252

Qy    241 CTGCCAAGGAGTACTGGATGTTCAAACCTCCGCAACTCCTCCAGCCAGCCACCGGGGAGG 300
      |||
Db   253 CTGCCAAGGAGTACTGGATGTTCAAACCTCCGCAACTCCTCCAGCCAGCCACCGGGGAGG 312

Qy    301 ACCCTGAGGGCTCAGACATCCTGAACTACTTTGAGAGCTACCTTGCCGTTGCCTCCACCG 360
      |||
Db   313 ACCCTGAGGGCTCAGACATCCTGAACTACTTTGAGAGCTACCTTGCCGTTGCCTCCACCG 372

Qy    361 TGCCCTCCATGTGTGCTGGTGGCCAACTTCCTGCTTGTCAACAGGGTTGCAGTCCACA 420
      |||
Db   373 TGCCCTCCATGTGTGCTGGTGGCCAACTTCCTGCTTGTCAACAGGGTTGCAGTCCACA 432

Qy    421 TCCGTGTCTTGGCCTCACTGACGGTCATCCTGGCCATCTTCATGGTGATAACTGCACTGG 480
      |||
Db   433 TCCGTGTCTTGGCCTCACTGACGGTCATCCTGGCCATCTTCATGGTGATAACTGCACTGG 492

Qy    481 TGAAGGTGGACACTTTCTCTGGACCCGTGGCTTTTTTGCGGTACCATGTGTGCATGG 540
      |||
Db   493 TGAAGGTGGACACTTCCTCCTGGACCCGTGGTTTTTTTGCGGTACCATGTGTGCATGG 552

Qy    541 TGATCCTCAGCGGTGCCTCCACTGTCTTTCAGCAGCAGCATCTACGGCATGACCGGCTCCT 600
      |||
Db   553 TGATCCTCAGCGGTGCCTCCACTGTCTTTCAGCAGCAGCATCTACGGCATGACCGGCTCCT 612

Qy    601 TTCCTATGAGGAACCTCCAGGCACCTGATATCAGGAGGAGCCATGGGCGGGACGGTCAGCG 660
      |||
Db   613 TTCCTATGAGGAACCTCCCAAGCAGTATATCAGGAGGAGCCATGGGCGGGACGGTCAGCG 672

Qy    661 CCGTGGCCTCATTGGTGGACTTGGCTGCATCCAGTGTGAGGAACAGCGCCCTGGCCT 720
      |||
Db   673 CCGTGGCCTCATTGGTGGACTTGGCTGCATCCAGTGTGAGGAACAGCGCCCTGGCCT 732

Qy    721 TCTTCCTGACGGCCACCATCTTCCTCGTGCTCTGCATGGGACTCTACCTGCTGCTGTCCA 780
      |||
Db   733 TCTTCCTGACGGCCACCATCTTCCTCGTGCTCTGCATGGGACTCTACCTGCTGCTGTCCA 792

Qy    781 GGCTGGAGTATGCCAGGTACTACATGAGGCCTGTTCTTGCGGCCCATGTGTTTTCTGGTG 840
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Art Unit: 1647

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Db      793  GGCTGGAGTATGCCAGGTACTACATGAGGCCTGTTCTTGCGGCCCATGTGTTTCTGGTG 852
Qy      841  AAGAGGAGCTTCCCCAGGACTCCCTCAGTGCCCCCTTCGGTGGCCTCCAGATTATTGATT 900
|||||
Db      853  AAGAGGAGCTTCCCCAGGACTCCCTCAGTGCCCCCTTCGGTGGCCTCCAGATTATTGATT 912
Qy      901  CCCACACACCCCCCTCTCCGCCCCATCCTGAAGAAGACGGCCAGCCTGGGCTTCTGTGTCA 960
|||||
Db      913  CCCACACACCCCCCTCTCCGCCCCATCCTGAAGAAGACGGCCAGCCTGGGCTTCTGTGTCA 972
Qy      961  CCTACGTCTTCTTCATCACCAGCCTCATCTACCCCGCCGTCTGCACCAACATCGAGTCCC 1020
|||||
Db      973  CCTACGTCTTCTTCATCACCAGCCTCATCTACCCCGCCGTCTGCACCAACATCGAGTCCC 1032
Qy      1021 TCAACAAGGGCTCGGGCTCACTGTGGACCACCAAGTTTTTCATCCCCCTCACTACCTTCC 1080
|||||
Db      1033 TCAACAAGGGCTCGGGCTCACTGTGGACCACCAAGTTTTTCATCCCCCTCACTACCTTCC 1092
Qy      1081 TCCTGTACAACTTTGCTGACCTATGTGGCCGGCAGCTCACCGCCTGGATCCAGGTGCCAG 1140
|||||
Db      1093 TCCTGTACAACTTTGCTGACCTATGTGGCCGGCAGCTCACCGCCTGGATCCAGGTGCCAG 1152
Qy      1141 GGCCCAATAGCAAGGCGCTCCCAGGGTTCGTGCTCCTCCGGACCTGCCTCATCCCCCTCT 1200
|||||
Db      1153 GGCCCAACAGCAAGGCGCTCCCAGGGTTCGTGCTCCTCCGGACCTGCCTCATCCCCCTCT 1212
Qy      1201 TCGTGCTCTGTAACTACCAGCCCCGCGTCCACCTGAAGACTGTGGTCTTCCAGTCCGATG 1260
|||||
Db      1213 TCGTGCTCTGTAACTACCAGCCCCGCGTCCACCTGAAGACTGTGGTCTTCCAGTCCGATG 1272
Qy      1261 TGTACCCCGCACTCCTCAGCTCCCTGCTGGGGCTCAGCAACGGCTACCTCAGCACCTGG 1320
|||||
Db      1273 TGTACCCCGCACTCCTCAGCTCCCTGCTGGGGCTCAGCAACGGCTACCTCAGCACCTGG 1332
Qy      1321 CCCTCCTCTACGGGCCTAAGATTGTGCCCAGGGAGCTGGCTGAGGCCACGGGAGTGGTGA 1380
|||||
Db      1333 CCCTCCTCTACGGGCCTAAGATTGTGCCCAGGGAGCTGGCTGAGGCCACGGGAGTGGTGA 1392
Qy      1381 TGTCTCTTTATGTGTGCTTGGGCTTAACACTGGGCTCAGCCTGCTCTACCCTCCTGGTGC 1440
|||||
Db      1393 TGTCTCTTTATGTGTGCTTGGGCTTAACACTGGGCTCAGCCTGCTCTACCCTCCTGGTGC 1452
Qy      1441 ACCTCATCTAGAAGGGAGGACACAAGGACATTGGTGCTTCAGAGCCTTTGAAGATGAGAA 1500
|||||
Db      1453 ACCTCATCTAGAAGGGAGGACACAAGGACATTGGTGCTTCAGAGCCTTTGAAGATGAGAA 1512
Qy      1501 GAGAGTGCAGGAGGGCTGGGGGCCATGGAGGAAAGGCCTAAAGTTTCACTTGGGGACAGA 1560
|||||
Db      1513 GAGAGTGCAGGAGGGCTGGGGGCCATGGAGGAAAGGCCTAAAGTTTCACTTGGGGACAGA 1572
Qy      1561 GAGCAGAGCACACTCGGGCCTCATCCCTCCCAAGATGCCAGTGAGCCACGTCCATGCCCA 1620
|||||
Db      1573 GAGCAGAGCACACTCGGGCCTCATCCCTCCCAAGATGCCAGTGAGCCACGTCCATGCCCA 1632
Qy      1621 TTCCGTGCAAGGCAGATATTCCAGTCATATTAACAGAACACTCCTGAGACAGTTGAAGAA 1680
|||||
Db      1633 TTCCGTGCAAGGCAGATATTCCAGTCATATTAACAGAACACTCCTGAGACAGTTGAAGAA 1692
Qy      1681 GAAATAGCACAAATCAGGGGTACTCCCTTCACAGCTGATGGTTAACATTCCACCTTCTTT 1740
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Art Unit: 1647

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      |||
Db      1693  GAAATAGCACAAATCAGGGGTACTCCCTTCACAGCTGATGGTTAACATTCCACCTTCTTT 1752
      |||
Qy      1741  CTAGCCCTTCAAAGATGCTGCCAGTGTTCGCCCTAGAGTTATTACAAAGCCAGTGCCAAA 1800
      |||
Db      1753  CTAGCCCTTCAAAGATGCTGCCAGTGTTCGCCCTAGAGTTATTACAAAGCCAGTGCCAAA 1812
      |||
Qy      1801  ACCCAGCCATGGGCTCTTTGCAACCTCCCAGCTGCGCTCATTCCAGCTGACAGCGAGATG 1860
      |||
Db      1813  ACCCAGCCATGGGCTCTTTGCAACCTCCCAGCTGCGCTCATTCCAGCTGACAGCGAGATG 1872
      |||
Qy      1861  CAAGCAAATGCTCAGTCTCCTTACCCTGAAGGGGTCTCCCTGGAATGGAAGTCCCCTGG 1920
      |||
Db      1873  CAAGCAAATGCTCAGTCTCCTTACCCTGAAGGGGTCTCCCTGGAATGGAAGTCCCCTGG 1932
      |||
Qy      1921  CATGGTCAGTCCCTCAGGCCCAAGACTCAAGTGTGCACAGACCCCTGTGTCTGTGGGTGA 1980
      |||
Db      1933  CATGGTCAGTCCCTCAGGCCCAAGACTCAAGTGTGCACAGACCCCTGTGTCTGTGGGTGA 1992
      |||
Qy      1981  ACAACTGCCCACTAACCAGACTGGAAAACCCAGAAAGATGGGCCTTCCATGAATGCTTCA 2040
      |||
Db      1993  ACAACTGCCCACTAACCAGACTGGAAAACCCAGAAAGATGGGCCTTCCATGAATGCTTCA 2052
      |||
Qy      2041  TTCCAGAGGGACCAGAGGGCCTCCCTGTGCAAGGGATCAAGCATGTCTGGCCTGGGTTT 2100
      |||
Db      2053  TTCCAGAGGGACCAGAGGGCCTCCCTGTGCAAGGGATCAAGCATGTCTGGCCTGGGTTT 2112
      |||
Qy      2101  CAAAAAAGAGGGATCCTCATGACCTGGTGGTCTATGGCCTGGGTCAAGATGAGGGTCTT 2160
      |||
Db      2113  CAAAAAAGAGGGATCCTCATGACCTGGTGGTCTATGGCCTGGGTCAAGATGAGGGTCTT 2172
      |||
Qy      2161  TCAGTGTTCCTGTTTACAACATGTCAAAGCCATTGGTTCAAGGGCGTAATAAAATACTTGC 2220
      |||
Db      2173  TCAGTGTTCCTGTTTACAACATGTCAAAGCCATTGGTTCAAGGGCGTAATAAAATACTTGC 2232
      |||
Qy      2221  GTATTCAAAA 2231
      |||
Db      2233  GTATTCAAAA 2243

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Conclusion:

No claims are allowed.

All claims are drawn to the same invention claimed in the application prior to the entry of the submission under 37 CFR 1.114 and could have been finally rejected on the grounds and art of record in the next Office action if they had been entered in the application prior to entry under 37 CFR 1.114. Accordingly, **THIS ACTION IS MADE FINAL** even though it is a first action after the filing of a request for continued

Art Unit: 1647

examination and the submission under 37 CFR 1.114. See MPEP § 706.07(b).

Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Deleted: ¶

Any inquiry concerning this communication or earlier communications from the examiner should be directed to SHULAMITH H. SHAFER whose telephone number is (571)272-3332. The examiner can normally be reached on Monday through Friday, 8 AM to 5 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Manjunath Rao, Ph.D. can be reached on 571-272-0939. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Lorraine Spector/ Ph.D.
Primary Examiner, Art Unit 1647

/S. H. S./
Examiner, Art Unit 1647